INDEX 'ADISCTI, ADISINSIGHT, ADISNEWS, AGRICOLA, ANABSTR, AQUASCI, BIOBUSINESS, BIOCOMMERCE, BIOSIS, BIOTECHABS, BIOTECHDS, BIOTECHNO, CABA, CANCERLIT, CAPLUS, CEABA-VTB, CEN, CIN, CONFSCI, CROPB, CROPU, DDFB, DDFU, DGENE, DRUGB, DRUGLAUNCH, DRUGMONOG2, ...' ENTERED AT 14:09:15 ON 02 JAN 2003

SEA ALPHA (W) 1, 6-FUCOSYLTRANSFERASE

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    FILE BIOCOMMERCE
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    FILE BIOSIS
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    FILE GENBANK
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    FILE LIFESCI
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533 FILE ESBIOBASE

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FILE FROSTI

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ANSWER 1 OF 11 CAPLUS COPYRIGHT 2003 ACS

2001:307991 CAPLUS ACCESSION NUMBER:

DOCUMENT NUMBER:

135:286390

TITLE:

Significance of .alpha.1-6 fucosylation in

hepatocellular carcinoma

AUTHOR(S):

Miyoshi, Eiji; Noda, Katsuhisa; Taniguchi, Naoyuki;

Sasaki, Yutaka; Hayashi, Norio

CORPORATE SOURCE:

Department of Biochemistry, Osaka University Graduate

School of Medicine, Suita, 565-0871, Japan

SOURCE:

Liver Cirrhosis (2001), 93-104. Editor(s): Okita,

Kiwamu. Springer-Verlag Tokyo: Tokyo, Japan.

CODEN: 69BFS7

DOCUMENT TYPE:

Conference; General Review

LANGUAGE:

English

A review, with 49 refs. .alpha.1-6

Fucosyltransferase (.alpha.1-6FucT) catalyzes the transfer of fucose from GDP-Fuc to N-linked type complex glycoproteins. Recently, serum .alpha.1-6 fucosylated .alpha.-fetoprotein (AFP) has been employed for a differential diagnosis of hepatocellular carcinoma (HCC) from liver cirrhosis. To det. the mol. basis of the fucosylated AFP in the serum of patients with HCC, we have succeeded in the purifn. and cDNA cloning of .alpha.1-6FucT from porcine brain and a gastric cancer cell line, resp. Their homol. is 92.2% at the nucleotide level and 95.7% at the amino acid level. No putative N-glycosylation sites were found in the predicted amino acid sequence. .alpha.1-6FucT was widely expressed in various rat tissues except normal liver. Expression of .alpha.1-6FucT in the liver was enhanced during hepatocarcinogenesis of LEC rats, which develop hereditary hepatitis and hepatomas. In human liver diseases, .alpha.1-6FucT was expressed in both HCCs and their surrounding tissues with chronic liver disease, but not in normal liver. Although serum AFP has been employed for an early diagnosis of patients with HCC, the mechanisms by which .alpha.1-6 fucosylation of AFP occurs in HCC seem to be not solely due to the up-regulation of .alpha.1-6FucT. Interestingly, when the .alpha.1-6FucT gene was transfected into Hep3B, a human hepatoma cell line, tumor formation in the liver of nude mice after splenic injection was dramatically suppressed. The mechanisms of the suppression were due to decreases in cell adhesion through aberrant glycosylation of .alpha.5.beta.1 integrin. In this review, we focus on the biol. significance of .alpha.1-6 fucosylation in HCC.

REFERENCE COUNT:

THERE ARE 49 CITED REFERENCES AVAILABLE FOR THIS 49 RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

ANSWER 2 OF 11 SCISEARCH COPYRIGHT 2003 ISI (R)

2000:459176 SCISEARCH ACCESSION NUMBER:

THE GENUINE ARTICLE: 323NE

TITLE:

High-mannose-type oligosaccharides from human placental

arylsulfatase A are core fucosylated as confirmed by MALDI

HojaLukowicz D (Reprint); Ciolczyk D; Bergquist J; AUTHOR:

Litynska A; Laidler P

CORPORATE SOURCE: JAGIELLONIAN UNIV, INST ZOOL, DEPT ANIM PHYSIOL, INGARDENA

6, PL-30060 KRAKOW, POLAND (Reprint); JAGIELLONIAN UNIV, COLL MEDICUM, INST MED BIOCHEM, PL-31034 KRAKOW, POLAND; UNIV GOTHENBURG, SAHLGRENS UNIV HOSP, INST CLIN NEUROSCI,

DEPT PSYCHIAT & NEUROCHEM, S-41380 MOLNDAL, SWEDEN

COUNTRY OF AUTHOR: POLAND; SWEDEN

GLYCOBIOLOGY, (JUN 2000) Vol. 10, No. 6, pp. 551-557. SOURCE:

Publisher: OXFORD UNIV PRESS, GREAT CLARENDON ST, OXFORD

OX2 6DP, ENGLAND. ISSN: 0959-6658.

DOCUMENT TYPE:

Article; Journal

FILE SEGMENT: LIFE LANGUAGE: English

REFERENCE COUNT:

ABSTRACT IS AVAILABLE IN THE ALL AND IALL FORMATS

Despite numerous studies on arylsulfatase A, the structure of its glycans is not well understood, It has been shown that the concentration of arylsulfatase A increases in the body fluids of patients with some forms of cancer, and the carbohydrate component of arylsulfatase A synthesized in tumor tissues and transformed cells undergoes increased sialylation, phosphorylation and sulfation. To understand the significance of any changes in the glycosylation of arylsulfatase A in cancer, it is important to know the structure of its carbohydrate component in normal tissue. In the present study we have analyzed carbohydrate moieties of human placental arylsylfatase A using sodium dodecyl sulfatepolyacrylamide gel electrophoresis (SDS-PAGE) followed by Western blotting on Immobilon P and on-blot deglycosylation using PNGase $\bar{\textbf{F}}$ for glycan release. Profiles of N-glycans were obtained by matrix-assisted laser desorption/ionization mass spectrometry (MALDI MS). Oligosaccharides were sequenced using specific exoglycosidases, and digestion products were analyzed by MALDI MS and the computer matching of the resulting masses with those derived from a sequence database. Fifty picomoles (6 mu g) of arylsulfatase A applied to the gel were sufficient to characterize its oligosaccharide content. The results indicated that human placental arylsulfatase A possesses only high-mannose-type oligosaccharides, of which almost half are core fucosylated. In addition, there was a minor species of high-mannose-type glycan bearing six mannose residues with a core fucose, This structure was not expected since high-mannose-type oligosaccharides basically have not been recognized as a substrate for the alpha 1,6-fucosyltransferase.

ANSWER 3 OF 11 CAPLUS COPYRIGHT 2003 ACS DUPLICATE 1

ACCESSION NUMBER: 2000:507631 CAPLUS

DOCUMENT NUMBER:

133:250111

TITLE:

Occurrence of GDP-L-fucose:.beta.-N-acetylglucosamine

(Fuc to Asn-linked GlcNAc) .alpha.1

,6-fucosyltransferases in

porcine, sheep, bovine, rabbit and chicken

tissues

AUTHOR(S):

Struppe, E.; Staudacher, E.

CORPORATE SOURCE:

Institut fur Chemie, Universitat fur Bodenkultur,

Vienna, A-1190, Austria

SOURCE:

Biochimica et Biophysica Acta (2000), 1475(3), 360-368

CODEN: BBACAQ; ISSN: 0006-3002

PUBLISHER:

Elsevier Science B.V.

DOCUMENT TYPE:

Journal

LANGUAGE:

English

Transgenic animals are a promising source of pharmaceutically-relevant proteins or as a source of organs for xenotransplantation. Beside other posttranslational modifications, glycosylation has been shown to be a crit. parameter for the correct function of several glycoproteins. To analyze the contribution of .alpha.1,6-fucosylation to N-glycan variability, the authors partly purified .alpha.

1,6-fucosyltransferase (.alpha.1,6-Fuc-T)

activities from various tissues (brain, lung, heart, liver) of agrıculturally-relevant animals (porcine, sheep, bovine, rabbit, chicken) and compared some of their biochem. properties. All tissues displayed .alpha.1,6-Fuc-T activity, although at different levels. No differences were obsd. in their stability against chems., temp. or time, whereas the activities were distinguishable by their pH-optima and their cation preferences. Similarities were found for tissues between species. Lung and heart enzymes showed a narrow pH-optimum around pH 6.0 and an enhanced activity in the presence of divalent cations. .alpha.1,6-Fuc-T activities in brain and liver were characterized by a broad pH-optimum from 5.5 to 8.0. Some activities of these tissues were

decreased by the addn. of EDTA, while others did not show any influence of EDTA or divalent cations. From the significant differences of the .alpha.1,6-Fuc-T activities in the tissues, it is possible to hypothesize the presence of more than one single .alpha.1,6-Fuc-T in mammalian tissues.

REFERENCE COUNT:

THERE ARE 32 CITED REFERENCES AVAILABLE FOR THIS 32 RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

DUPLICATE 2

ANSWER 4 OF 11 CAPLUS COPYRIGHT 2003 ACS

1999:746555 CAPLUS

ACCESSION NUMBER: DOCUMENT NUMBER:

132:119026

TITLE:

The .alpha.1-6-

fucosyltransferase gene and its biological

significance

AUTHOR(S):

Miyoshi, E.; Noda, K.; Yamaguchi, Y.; Inoue, S.; Ikeda, Y.; Wang, W.; Ko, J. H.; Uozumi, N.; Li, W.;

Taniguchi, N.

CORPORATE SOURCE:

Department of Biochemistry, Osaka University Medical

School, Suita, Osaka, Japan

SOURCE:

Biochimica et Biophysica Acta (1999), 1473(1), 9-20

CODEN: BBACAO; ISSN: 0006-3002

PUBLISHER: DOCUMENT TYPE: Elsevier Science B.V. Journal; General Review

English

LANGUAGE:

A review with 94 refs. GDP-L-fucose: N-acetyl-.beta.-D-glucosaminide .

alpha.1-6-fucosyltransferase

(.alpha.1-6FucT) catalyzes the transfer of fucose from GDP-fucose to N-linked type complex glycoproteins. This enzyme was purified from a human fibroblast cell line, porcine brain, a human gastric cancer cell line and human blood platelets. cDNA cloning of porcine and human .alpha.1-6FucT was performed from a porcine brain and gastric cancer cell cDNA libraries, resp. Their homol. is 92.2% at the nucleotide level and 95.7% at the amino acid level. No putative N-glycosylation sites were found in the predicted amino acid sequence. No homol. to other fucosyltransferases such as .alpha.1-2FucT, .alpha.1-3FucT and .alpha.1-4FucT was found except for a region consisting of nine amino acids. The .alpha.1-6FucT gene is located at chromosome 14q24.3, which is also a different location from other fucosyltransferases reported to date. The .alpha.1-6FucT gene is the oldest gene family in the phylogenic trees among the nine cloned fucosyltransferase genes.

.alpha.1-6FucT is widely expressed in various rat tissues and the expression of .alpha.1-6FucT in the liver is enhanced during hepatocarcinogenesis of LEC rats which develop hereditary hepatitis and hepatomas. In cases of human liver diseases, .alpha.1-6FucT is expressed in both hepatoma tissues and their surrounding tissues with chronic liver disease, but not in the case of normal liver. Serum .alpha.1-6fucosylated .alpha.-fetoprotein (AFP) has been employed for an early diagnosis of patients with hepatoma. The mechanisms by which .alpha.1-6 fucosylation of AFP occurs in the hepatoma is not due to the up-regulation of .alpha.1-6FucT alone. Interestingly, when the .alpha.1-6FucT gene is transfected into Hep3B, a human hepatoma cell line, tumor formation in the liver of nude mice after splenic injection is dramatically suppressed. This review focuses on .alpha.1-6FucT and summarizes its properties, gene

expression and biol. significance.

REFERENCE COUNT:

94 THERE ARE 94 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

ANSWER 5 OF 11 CAPLUS COPYRIGHT 2003 ACS

DUPLICATE 3

ACCESSION NUMBER:

1998:680686 CAPLUS

DOCUMENT NUMBER:

130:79497

TITLE:

Gene expression of .alpha.1-6 fucosyltransferase in human

hepatoma tissues: a possible implication for increased

fucosylation of .alpha.-fetoprotein

Noda, Katsuhisa; Miyoshi, Eiji; Uozumi, Naofumi; AUTHOR(S):

> Yanaqidani, Shusaku; Ikeda, Yoshitaka; Gao, Cong-Xiao; Suzuki, Kunio; Yoshihara, Harumasa; Yoshikawa, Masumi; Kawano, Kiyoshi; Hayashi, Norio; Hori, Masatsuqu;

Taniguchi, Naoyuki CORPORATE SOURCE:

Department of Biochemistry, Osaka University Medical

School, Osaka, 565-0871, Japan

Hepatology (Philadelphia) (1998), 28(4), 944-952 SOURCE:

CODEN: HPTLD9; ISSN: 0270-9139

W. B. Saunders Co. PUBLISHER:

Journal DOCUMENT TYPE: English LANGUAGE:

The .alpha.1-6 fucosylated .alpha.-fetoprotein (AFP) present in serum of patients with hepatocellular carcinoma (HCC) has been employed for the differential clin. diagnosis of HCC from chronic liver diseases. The mol. mechanism by which this alteration occurs, however, remains largely unknown. To address this issue, the authors previously purified GDP-L-Fuc: N-acetyl-.beta.-D-glucosaminide .alpha.1-

6 fucosyltransferase (.alpha.1-6 FucT), an enzyme involved in the .alpha.1-6 fucosylation of N-qlycans from porcine brain, as well as from a human gastric cancer cell line, and cloned their genes. In this study, levels of .alpha.1-6 FucT mRNA expression and the activity of this enzyme for 12 human HCC tissues were examd. and compared with that in surrounding tissues and normal livers. The mean for .alpha.1-6 FucT activity was 78 pmol/h/mg in normal control liver, 202 pmol/h/mg in adjacent uninvolved liver tissues (chronic hepatitis: 181 pmol/h/mg; liver cirrhosis: 233 pmol/h/mg), and 195 pmol/h/mg in HCC tissues. The mRNA expression of .alpha.1-6 FucT was also enhanced in proportion to enzymic activity except for a few cases, suggesting that .alpha.1-6 FucT expression is increased in chronic liver diseases, esp. liver cirrhosis. Transfection of .alpha.1-6 FucT gene into cultured rat hepatocytes markedly increased .alpha.1-6 FucT activity and led to an increase in lens culinaris agglutinin (LCA) binding proteins in both cell lysates and condition media. When the .alpha.1-6 FucT gene was transfected into a human HCC cell line, Hep3B, which originally showed low levels of .alpha.1-6 FucT expression, .alpha.1-6-fucosylated AFP was dramatically increased in the condition media. Collectively, these results suggest that the enhancement of .alpha.1-6 FucT expression increased the fucosylation of several proteins, including AFP, and that

caused by up-regulation of the .alpha.1-6 FucT gene expression. THERE ARE 32 CITED REFERENCES AVAILABLE FOR THIS REFERENCE COUNT: 32 RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

the level of .alpha.1-6-fucosylated AFP in patients with HCC was in part

ANSWER 6 OF 11 CAPLUS COPYRIGHT 2003 ACS DUPLICATE 4

ACCESSION NUMBER: 1998:95524 CAPLUS

DOCUMENT NUMBER: 128:165712

TITLE: High expression of .alpha.-1-

6 fucosyltransferase during rat

hepatocarcinogenesis

Noda, Katsuhisa; Mitoshi, Eiji; Uozumi, Naofumi; Gao, AUTHOR(S):

Cong-Xiao; Suzuki, Keiichiro; Hayashi, Norio; Hori,

Masatsugu; Taniquchi, Naoyuki

CORPORATE SOURCE: Department of Biochemistry, Osaka University Medical

School, Osaka, 565, Japan

SOURCE: International Journal of Cancer (1998), 75(3), 444-450

CODEN: IJCNAW; ISSN: 0020-7136

PUBLISHER: Wiley-Liss, Inc.

DOCUMENT TYPE: Journal

LANGUAGE: English

.alpha.-1-6 Fucosylated .alpha.-fetoprotein (AFP) is known to be elevated in patients with primary hepatoma and has been suggested as being useful as an early indicator and predictor of the poor prognosis for hepatoma. Although GDP-L-fucosyl-N-acetyl-.beta.-D-glucosaminide .alpha.-

1-6 fucosyltransferase (.alpha.-1-6 FucT), is the key enzyme involved in .alpha.-1-6 fucosylation of AFP, when and how the expression of .alpha.-1-6 FucT is enhanced during hepatocarcinogenesis is unknown. Recently, we established a convenient assay method for this enzyme and were successful in the purifn. and cDNA cloning of .alpha.-1-6 FucT from human gastric cancer, as well as from porcine brain. In the present study, levels of .alpha.-1-6 FucT activity and mRNA expression have been detd. during hepatocarcinogenesis in LEC rats which spontaneously develop hereditary hepatitis and hepatoma. The fetal liver contained the highest enzymic activity, which tended to increase in inverse proportion to gestation. The enzymic activity was significantly increased in hepatoma tissues as compared with uninvolved adjacent tissues. Northern-blot anal. revealed high expression of .alpha.-1-6 FucT mRNA in hepatoma tissues, whereas the expression was fairly low in normal, hepatitis and uninvolved adjacent liver tissues. While the fetal liver had the highest enzymic activity, the expression of .alpha.-1-6 FucT mRNA was low, suggesting that another .alpha.-1-6 FucT is induced in fetal liver or that post-translational modification occurs. High expression of .alpha.-1-6 FucT was also obsd. in 3'-MeDAB-induced rat hepatomas and some rat hepatoma cell lines. .alpha.-1-6 FucT was strongly enhanced from an early stage of hepatocarcinogenesis and was maintained at a high level in rat hepatomas.

L5 ANSWER 7 OF 11 CAPLUS COPYRIGHT 2003 ACS DUPLICATE 5

ACCESSION NUMBER: 1997:652209 CAPLUS

DOCUMENT NUMBER:

127:329624

TITLE:

Expression of .alpha.1-6

fucosyltransferase in rat tissues and human

cancer cell lines

AUTHOR(S):

SOURCE:

Miyoshi, Eiji; Uozumi, Naofumi; Noda, Katsuhisa; Hayashi, Norio; Hori, Masatsugu; Taniguchi, Naoyuki Department of Biochemistry, Osaka University Medical

CORPORATE SOURCE:

School, Suita, 565, Japan International Journal of Cancer (1997), 72(6),

1117-1121

CODEN: IJCNAW; ISSN: 0020-7136

PUBLISHER: Wiley-Liss
DOCUMENT TYPE: Journal
LANGUAGE: English

AB GDP-L-Fuc: N-acetyl-.beta.-D-glucosaminide .alpha.1-

6 fucosyltransferase (.alpha.1-6FucT) catalyzed the transfer of a fucosyl residue from GDP-fucose to the asparagine-linkage GlcNAc residue of complex N-glycans via .alpha.1-6 linkage. These oligosaccharide structures are essential for the attachment of polysialic acid to the neural-cell-adhesion mol., and its levels are useful for the differential diagnosis of hepatocellular carcinomas with respect to the microheterogeneity of .alpha.-fetoprotein. The authors have been successful in the purifn. of cDNA cloning of .alpha.1-6FucT from porcine brain and from a human gastric-cancer cell line. In the present study, mRNA expression of .alpha.1-6FucT in various rat tissues and human cancer cell lines was examd., along with the expression of .alpha.1-6FucT mRNA and the induction by treatment with several cytokines. Northern-blot anal. indicated high expression levels of .alpha.1-6FucT in brain and gastrointestinal-tract tissues of normal rats, as well as for a no. of lung-cancer, gastric-cancer and colon-cancer lines. Although various cytokines did not induce .alpha.1-6FucT mRNA, differentiation of a tumor cell enhanced the mRNA by 2- to 3-fold. These results may provide new insight into studies on .alpha.1-6FucT in terms of carcinogenesis or differentiation.

5 ANSWER 8 OF 11 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC.

ACCESSION NUMBER: 1998:58067 BIOSIS DOCUMENT NUMBER: PREV199800058067

TITLE: Purification and cDNA cloning of porcine

brain GDP-L-Fuc: N-acetyl-beta-D-glucosaminide: alpha1-

6fucosyltransferase.

AUTHOR(S): Uozumi, N. (1); Yanaqidani, S.; Miyoshi, E. (1); Ihara, Y.

(1); Sakuma, T. (1); Kang, R. (1); Gao, C.-X. (1); Noda, K. (1); Teshima, T.; Fujii, S.; Shiba, T.; Taniguchi, N. (1)

CORPORATE SOURCE: (1) Dep. Biochem., Osaka Univ. Med. Sch., 2-2 Yamadaoka,

Suita, Osaka 565 Japan

SOURCE: Glycoconjugate Journal, (1997) Vol. 14, No. 6, pp. 762.

Meeting Info.: International Symposium on

Glycosyltransferases and Cellular Communications Osaka,

Japan March 26-28, 1997

ISSN: 0282-0080.

DOCUMENT TYPE: LANGUAGE:

CORPORATE SOURCE:

PUBLISHER:

Conference English

L5 ANSWER 9 OF 11 CAPLUS COPYRIGHT 2003 ACS DUPLICATE 6

ACCESSION NUMBER: 1997:261195 CAPLUS

DOCUMENT NUMBER: 126:327408

TITLE: Purification and cDNA cloning of

GDP-L-Fuc: N-acetyl-.beta.-D-glucosaminide:.

alpha.1-6

fucosyltransferase (.alpha.1-6 FucT) from

human gastric cancer MKN45 cells

AUTHOR(S): Yanagidani, Shusaku; Uozumi, Naofumi; Ihara, Yoshito;

Miyoshi, Eiji; Yamaguchi, Nozomi; Taniguchi, Naoyuki Department of Biochemistry, Osaka University Medical

School, Osaka, 565, Japan

SOURCE: Journal of Biochemistry (Tokyo) (1997), 121(3),

626-632

CODEN: JOBIAO; ISSN: 0021-924X Japanese Biochemical Society

DOCUMENT TYPE: Journal LANGUAGE: English

AB GDP-L-Fuc:N-acetyl-.beta.-D-glucosaminide:.alpha.1-

6 fucosyltransferase (.alpha.1-6 FucT), which catalyzes the transfer of fucose from GDP-Fuc to N-linked type complex glycopeptides, was purified from a culture supernatant of human gastric cancer cell line MKN45. The purifn. procedures included chromatogs. on Q-Sepharose Fast Flow, synthetic GDP-hexanolamine-Sepharose, and GnGn-bi-Asn-Sepharose columns. SDS-PAGE of the purified enzyme gave a major band corresponding to an apparent mol. mass of 60 kDa. The enzyme was recovered in a 12% final yield with an approx. 4,600-fold increase in specific activity. The pH optimum was 7.5, and the enzyme was fully active in the presence of 5 $\overline{\text{mM}}$ EDTA and did not require divalent cations, Mg2+ and Ca2+. Oligonucleotide primers designed from partial amino acid sequences were used to amplify and clone .alpha.1-6 FucT cDNA from a cDNA library of MKN45 cells. The cDNA encodes 575 amino acids in length, and contains the predicted N-terminal and internal amino acid sequences derived on lysyl endopeptidase digestion. The homol. to porcine brain .alpha.1-6 FucT is 92.2% at the nucleotide level and 95.7% at the amino acid level. No putative N-glycosylation sites were found in the predicted amino acid sequence of the human MKN45 cell enzyme or that of porcine brain. Thus, the enzyme is distinct from other fucosyltransferases which catalyze .alpha.1-2, .alpha.1-3, and .alpha.1-4 fucose addn.

L5 ANSWER 10 OF 11 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC.

ACCESSION NUMBER: 1997:233962 BIOSIS DOCUMENT NUMBER: PREV199799533165

TITLE: Alpha-1-6

Fucosyltransferase: Purification, cDNA

cloning and expression during hepatocarcinogenesis.

AUTHOR(S): Miyoshi, E.; Uozumi, N.; Noda, K.; Taniguchi, N.

CORPORATE SOURCE: Osaka Univ. Med. Sch., Osaka Japan

SOURCE:

Proceedings of the American Association for Cancer Research

Annual Meeting, (1997) Vol. 38, No. 0, pp. 561.

Meeting Info.: Eighty-eighth Annual Meeting of the American Association for Cancer Research San Diego, California, USA

April 12-16, 1997 ISSN: 0197-016X. Conference; Abstract

DOCUMENT TYPE:

LANGUAGE: English

ANSWER 11 OF 11 BIOSIS COPYRIGHT 2003 BIOLOGICAL ABSTRACTS INC.

ACCESSION NUMBER: DOCUMENT NUMBER:

1996:574476 BIOSIS PREV199799289157

TITLE:

Purification and cDNA cloning of porcine

brain GDP-L-Fuc: N-acetyl-beta-D-glucosaminide alpha-1

fwdarw 6fucosyltransferase.

AUTHOR (S):

Uozumi, Naofumi; Yanagidani, Shusaku; Miyoshi, Eiji; Ihara,

Yoshito; Sakuma, Takahiko; Gao, Cong-Xiao; Teshima,

Tadashi; Fujii, Shigeru; Shiba, Tetsuo; Taniquchi, Naoyuki

(1)

CORPORATE SOURCE:

(1) Dep. Biochemistry, Osaka Univ., Med. Sch., 2-2

Yamadaoka, Suita, Osaka 565 Japan

SOURCE:

Journal of Biological Chemistry, (1996) Vol. 271, No. 44,

pp. 27810-27817.
ISSN: 0021-9258.

DOCUMENT TYPE: LANGUAGE: Article English

GDP-L-Fuc:N-acetyl-beta-D-glucosaminide alpha-1 fwdarw 6fucosyl-transferase (alpha-1-6FucT; EC 2.4.1.68), which catalyzes the transfer of fucose from GDP-Fuc to N-linked type complex glycopeptides, was purified from a Triton X-100 extract of porcine brain microsomes. The purification procedures included sequential affinity chromatographies on GlcNAc-beta-1-2Man-alpha-1-6 (GlcNAc-beta-1-2Man-alpha-1-2) Man-beta-1-4GlcNAc-beta-1-4GlcNAc-Asn-Sepharose 4B and synthetic GDP-hexanolamine-Sepharose 4B columns. The enzyme was recovered in a 12% final yield with a 440,000-fold increase in specific activity. SDS-polyacrylamide gel electrophoresis of the purified enzyme gave a major band corresponding to an apparent molecular mass of 58 kDa. The alpha-1-6FucT has 575 amino acids and no putative N-glycosylation sites. The cDNA was cloned in to pSVK3 and was then transiently transfected into COS-1 cells. alpha-1-6FucT activity was found to be high in the transfected cells, as compared with non- or mock-transfected cells. Northern blotting analyses of rat adult tissues showed that alpha-1-6FucT was highly expressed in brain. No sequence homology was found with other previously cloned fucosyltransferases, but the enzyme appears to be a type II transmembrane protein like the other glycosyltransferases.

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	Terms	Documents
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	Search History	

DATE: Thursday, January 02, 2003 Printable Copy Create Case

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•	T,PGPB,JPAB,EPAB,DWPI; PLUR=YES; OP=ADJ	r	result set
<u>L3</u>	L2 same (isolat\$n, purif\$n, charact\$n)	0	<u>L3</u>
<u>L2</u>	L1 same (pig or porc\$n)	29	<u>L2</u>
<u>L1</u>	fucosyltransferase	424	<u>L1</u>

END OF SEARCH HISTORY

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L2: Entry 1 of 29

File: PGPB

Nov 28, 2002

PGPUB-DOCUMENT-NUMBER: 20020177551

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020177551 A1

TITLE: Compositions and methods for treatment of neoplastic disease

PUBLICATION-DATE: November 28, 2002

INVENTOR - INFORMATION:

NAME

CITY

STATE

COUNTRY

RULE-47

Terman, David S.

Pebble Beach

CA

US

US-CL-CURRENT: 514/12; 435/325, 530/350

Full Title Citation Front Review Classification Date Reference Sequences Attachments Claims KintC Draw Desc Image

L2: Entry 2 of 29

File: PGPB

Nov 7, 2002

PGPUB-DOCUMENT-NUMBER: 20020164749

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020164749 A1

TITLE: Alpha1,3-fucosyltransferase

PUBLICATION-DATE: November 7, 2002

INVENTOR - INFORMATION:

NAME

CITY

STATE

COUNTRY

RULE-47

Taylor, Diane E.

Ge, Zhongming

Edmonton Edmonton

CA CA

US-CL-CURRENT: 435/193; 435/320.1, 435/325, 435/69.1, 435/84, 536/23.2, 536/53

Full Title Citation Front Review Classification Date Reference Sequences Attachments Claims Mill Draw Desc. Image

☐ 3. Document ID: US 20020133836 A1

L2: Entry 3 of 29

File: PGPB

Sep 19, 2002

PGPUB-DOCUMENT-NUMBER: 20020133836

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020133836 A1

TITLE: Methods to identify swine genetically resistant to F18 E. coli associated diseases

PUBLICATION-DATE: September 19, 2002

INVENTOR-INFORMATION:

NAME

CITY

STATE

COUNTRY

RULE-47

Bosworth, Brad T.

Littleton

NC

US

Vogeli, Peter

Zurich

CH

US-CL-CURRENT: 800/17; 435/193, 435/6

Full Title Citation Front Review Classification Date Reference Sequences Attachments

KMMC | Drawn Desc | Image |

□ 4. Document ID: US 20020129395 A1

L2: Entry 4 of 29

File: PGPB

Sep 12, 2002

PGPUB-DOCUMENT-NUMBER: 20020129395

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020129395 A1

TITLE: Compositions to identify swine genetically resistant to F18 E. coli

associated diseases

PUBLICATION-DATE: September 12, 2002

INVENTOR-INFORMATION:

NAME

CITY

STATE

COUNTRY

RULE-47

Bosworth, Brad T.

Vogeli, Peter

Zurich

Littleton

NC

US CH

US-CL-CURRENT: 800/17; 435/193, 435/6, 536/23.2

Full Title Citation Front Review Classification Date Reference Sequences Attachments

KMC Draw Desc Image

L2: Entry 5 of 29

File: PGPB

Jun 27, 2002

PGPUB-DOCUMENT-NUMBER: 20020081694

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020081694 A1

TITLE: Alpha 1-6 fucosyltransferase

PUBLICATION-DATE: June 27, 2002

INVENTOR - INFORMATION:

NAME CITY STATE COUNTRY RULE-47

Taniguchi, Naoyuki Toyonaka-shi JP
Uozumi, Naofumi Kobe-shi JP
Shiba, Tetsuo Toyonaka-shi JP
Yanagidani, Shusaku Ohtsu-shi JP

US-CL-CURRENT: 435/193; 435/101, 435/320.1, 435/325, 435/69.1

Folto: Title | Citation | Front | Review | Classification | Date | Reference | Sequences | Attachments

FindC - Draw Dezi - Image

☐ 6. Document ID: US 20020068347 A1

L2: Entry 6 of 29

File: PGPB

Jun 6, 2002

PGPUB-DOCUMENT-NUMBER: 20020068347

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020068347 A1

TITLE: Nucleic acids encoding alpha-1,3 fucosyltransferases and expression systems

for making and expressing them

PUBLICATION-DATE: June 6, 2002

INVENTOR-INFORMATION:

NAME

CITY

STATE

COUNTRY

RULE-47

Taylor, Diane E.

Ge, Zhongming

Edmonton Edmonton CA CA

US-CL-CURRENT: 435/193; 435/325, 435/6, 435/69.1, 435/7.92, 530/389.1, 536/23.2

Full Title Citation Front Review Classification Date Reference Sequences Attachments

1960C Praw Desc Image

☐ 7. Document ID: US 20020037570 A1

L2: Entry 7 of 29

File: PGPB

Mar 28, 2002

PGPUB-DOCUMENT-NUMBER: 20020037570

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020037570 A1

TITLE: Alpha 1,2-fucosyltransferase

PUBLICATION-DATE: March 28, 2002

INVENTOR-INFORMATION:

NAME

CITY

STATE

COUNTRY

RULE-47

Taylor, Diane

Edmonton

CA

Wang, Ge

Edmonton

CA

Palcic, Monica

Edmonton

CA

US-CL-CURRENT: 435/193; 536/23.2

Full Title Citation Front Review Classification Date Reference Sequences Attachinents

KindC - Draw Desc - Image

■ 8. Document ID: US 20020031494 A1

L2: Entry 8 of 29

File: PGPB

Mar 14, 2002

PGPUB-DOCUMENT-NUMBER: 20020031494

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020031494 A1

TITLE: NUCLEIC ACIDS FOR REDUCING CARBOHYDRATE EPITOPES

PUBLICATION-DATE: March 14, 2002

INVENTOR - INFORMATION:

NAME

CITY

STATE COUNTRY

RULE-47

SANDRIN, MAURO SERGIO

BRUNSWICK

ΑU

MCKENZIE, IAN CAMPBELL FARQUHAR

BRUNSWICK

ΑU

US-CL-CURRENT: 424/93.2; 424/93.21, 435/320.1, 435/325, 435/455, 514/44, 536/23.2

Full Title Citation Front Review Classification Date Reference Sequences Attachments

KMC Draw Desc Image

□ 9. Document ID: US 20020028205 A1

L2: Entry 9 of 29

File: PGPB

Mar 7, 2002

PGPUB-DOCUMENT-NUMBER: 20020028205

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020028205 A1

TITLE: ANTIGENIC FUSIONPROTEIN CARRYING GALALPHA 1,3GAL EPITOPES

PUBLICATION-DATE: March 7, 2002

INVENTOR-INFORMATION:

NAME

CITY

STATE

COUNTRY SE

RULE-47

HOLGERSSON, JAN

LIU, JINING

HUDDINGE HUDDINGE

SE

US-CL-CURRENT: 424/184.1; 424/178.1, 435/320.1, 530/350, 530/387.1, 530/391.1,

536/23.1

Full Title Otation Front Review Classification Date Reference Sequences Attachments

MillO Draw Desc Image

☐ 10. Document ID: US 20020013957 A1

L2: Entry 10 of 29

File: PGPB

Jan 31, 2002

PGPUB-DOCUMENT~NUMBER: 20020013957

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020013957 A1

TITLE: Method of cloning porcine animals

PUBLICATION-DATE: January 31, 2002

INVENTOR-INFORMATION:

NAME

CITY

STATE COUNTRY

US

US

RULE-47

Damiani, Philip Betthauser, Jeffrey M.

Forsberg, Erik J.

Bishop, Michael D.

Windsor Oregon

Spencer

MA US WI US

WI Rio WI

US-CL-CURRENT: 800/24; 800/17

Full Title Citation Front Review Classification Cate Reference Sequences Attachiments

FindC - Draw Desc - Invage

☐ 11. Document ID: US 20010055584 A1

L2: Entry 11 of 29

File: PGPB

Dec 27, 2001

PGPUB-DOCUMENT-NUMBER: 20010055584

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20010055584 A1

TITLE: IMPROVED NUCLEIC ACIDS ENCODING A CHIMERIC GLYCOSYLTRANSFERASE

PUBLICATION-DATE: December 27, 2001

INVENTOR-INFORMATION:

NAME

CITY

STATE COUNTRY RULE-47

MCKENZIE, IAN FARQUHAR CAMPBELL

BRUNSWICK

ΑU

SANDRIN, MAURO SERGIO

BRUNSWICK

ΑU

US-CL-CURRENT: 424/93.2; 424/93.21, 435/320.1, 435/325, 435/455, 514/44, 536/23.1,

536/23.4

KMC Draw Desc Image

12. Document ID: US 6455037 B1

L2: Entry 12 of 29

File: USPT

Sep 24, 2002

US-PAT-NO: 6455037

DOCUMENT-IDENTIFIER: US 6455037 B1

TITLE: Cells expressing an .alpha.gala nucleic acid and methods of

xenotransplantation

Full Title Citation Front Review Classification Date Reference Sequences Attachments

Full Title Citation Front Review Classification Date Reference Sequences Attachments

KMMC Draw Desc Image

L2: Entry 13 of 29

File: USPT

Sep 3, 2002

US-PAT-NO: 6444655

DOCUMENT-IDENTIFIER: US 6444655 B1

TITLE: Galactopyranosides and their use

Full Title Citation Front Review Classification Date Reference Sequences Attachments

khmic Draw Desc Image

1 14. Document ID: US 6399758 B1

L2: Entry 14 of 29

File: USPT

Jun 4, 2002

US-PAT-NO: 6399758

DOCUMENT-IDENTIFIER: US 6399758 B1

TITLE: Nucleic acids for reducing carbohydrate epitopes

Full Title Citation Front Review Classification Date Reference Sequences Attachments FindE - Drawn Den - Innage 1 15. Document ID: US 6399337 B1 L2: Entry 15 of 29 File: USPT Jun 4, 2002 US-PAT-NO: 6399337 DOCUMENT-IDENTIFIER: US 6399337 B1 TITLE: .alpha.1,3-fucosyltransferase Full Title Citation Front Review Classification Date Reference Sequences Attachments PMC Draw Desc Image ☐ 16. Document ID: US 6291219 B1 L2: Entry 16 of 29 File: USPT Sep 18, 2001 US-PAT-NO: 6291219 DOCUMENT-IDENTIFIER: US 6291219 B1 TITLE: .alpha.1-6 fucosyltransferase Full Title Citation Front Review Classification Date Reference Sequences Attachments MadC Draw Desc Image ☐ 17. Document ID: US 6238894 B1 File: USPT L2: Entry 17 of 29 May 29, 2001 US-PAT-NO: 6238894 DOCUMENT-IDENTIFIER: US 6238894 B1 TITLE: .alpha.1,2 fucosyltransferase Full Title Citation Front Review Classification Date Reference Sequences Attachinents L2: Entry 18 of 29 Mar 20, 2001 File: USPT US-PAT-NO: 6204431 DOCUMENT-IDENTIFIER: US 6204431 B1 TITLE: Transgenic non-human mammals expressing heterologous glycosyltransferase DNA sequences produce oligosaccharides and glycoproteins in their milk Title Citation Front Review Classification Date Reference Sequences Attachments PMC | Draw Desc | Image

☐ 19. Document ID: US 6166288 A

L2: Entry 19 of 29

File: USPT

Dec 26, 2000

US-PAT-NO: 6166288

DOCUMENT-IDENTIFIER: US 6166288 A

TITLE: Method of producing transgenic animals for xenotransplantation expressing both an enzyme masking or reducing the level of the gal epitope and a complement inhibitor

Full Title Citation Front Remem Classification Date Reference Sequences Attachments

KNMC Draw Desc Image

□ 20. Document ID: US 6054304 A

L2: Entry 20 of 29

File: USPT

Apr 25, 2000

US-PAT-NO: 6054304

DOCUMENT-IDENTIFIER: US 6054304 A

TITLE: .alpha.1-6 fucosyltransferase

Full Title Citation Front Review Classification Date Reference Sequences Attachments

10x10 Pravi Desc Image

L2: Entry 21 of 29

File: USPT

Apr 6, 1999

US-PAT-NO: 5892070

DOCUMENT-IDENTIFIER: US 5892070 A

TITLE: Transgenic non-human mammals producing oligosaccharides and glycoconjugates

Full Title Citation Front Review Classification Date Reference Sequences Attachments

ROMC Draw Desc Image

L2: Entry 22 of 29

File: USPT

Apr 6, 1999

US-PAT-NO: 5891698

DOCUMENT-IDENTIFIER: US 5891698 A

TITLE: Oligosaccharides and glycoproteins produced in milk of transgenic non-human

mammals

Full Title Citation Front Review Classification Date Reference Sequences Attachments

PMIC Draw Desc Image

L2: Entry 23 of 29

File: USPT

Jan 12, 1999

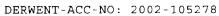
US-PAT-NO: 5858752

DOCUMENT-IDENTIFIER: US 5858752 A

TITLE: Fucosyltransferase genes and uses thereof

L2: Entry 28 of 29

File: DWPI Dec 18, 2001



DERWENT-WEEK: 200214

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TITLE: Producing pig organs with decreased immunogenicity due to modulation of sialyltransferase or a <u>fucosyltransferase</u> expression, useful as a source of donor organs for transplant into humans

Full : Title | Citation | Front | Remem | Classification | Date | Reference | Sequences | Attachments

PaulC - Draw Desc - Image

____ 29. Document ID: WO 9727303 A1 US 20020081694 A1 JP 09201191 A EP 816503 A1 JP 10004959 A JP 10004969 A JP 10084975 A US 6054304 A US 6291219 B1

L2: Entry 29 of 29

File: DWPI

Jul 31, 1997

DERWENT-ACC-NO: 1997-393690

DERWENT-WEEK: 200245

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TITLE: Human or pig alpha 1-6 fucosyl:transferase and DNA encoding it - for synthesis and modification of sugar chains and used as an antigen for production of diagnostic antibodies

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	Ge	enerate Colle	ection	Print		
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